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=> s l14(5a) (variant or muta? or substitution)

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PROCESSING COMPLETED FOR L15

L16 42 DUP REM L15 (36 DUPLICATES REMOVED)

=> d 1-10

- L16 ANSWER 1 OF 42 SCISEARCH COPYRIGHT (c) 2006 The Thomson Corporation on STN
- AN 2006:427530 SCISEARCH
- GA The Genuine Article (R) Number: 0240W
- TI Kinetic analysis of organophosphorus nerve agent hydrolysis by wildtype and mutant human paraoxonase 1
- AU Yeung D (Reprint); Cerasoli D; Lenz D
- CS USA, Med Res Inst Chem Def, Aberdeen Proving Ground, MD 21010 USA; Univ Maryland, Baltimore, MD 21201 USA
- CYA USA
- SO FASEB JOURNAL, (6 MAR 2006) Vol. 20, No. 4, Part 1, pp. A480-A480. ISSN: 0892-6638.
- PB FEDERATION AMER SOC EXP BIOL, 9650 ROCKVILLE PIKE, BETHESDA, MD 20814-3998 USA.
- DT Conference; Journal
- LA English
- REC Reference Count: 0
- ED Entered STN: 4 May 2006 Last Updated on STN: 4 May 2006
- L16 ANSWER 2 OF 42 MEDLINE on STN
- AN 2006259093 MEDLINE
- DN PubMed ID: 16683402
- TI Autopsy and postmortem examination case study on genetic risk factors for cardiac death: polymorphisms of endothelial nitric oxide synthase gene Glu298Asp variant and T-786C mutation, human

paraoxonase 1 (PON1) gene and alpha2beta-adrenergic receptor gene.

- AU Ameno Kiyoshi; Ameno Setsuko; Kinoshita Hiroshi; Jamal Mostofa; Wang Weihuan; Kumihashi Mitsuru; Uekita Ikuo; Ijiri Iwao
- CS Department of Forensic Medicine, Faculty of Medicine, Kagawa University, Japan.. amenok@kms.ac.jp
- SO Vojnosanitetski pregled. Military-medical and pharmaceutical review, (2006 Apr) Vol. 63, No. 4, pp. 357-61; discussion 362-3.

 Journal code: 21530700R. ISSN: 0042-8450.
- CY Serbia and Montenegro
- DT Journal; Article; (JOURNAL ARTICLE)
- LA English
- FS Priority Journals
- EM 200605
- ED Entered STN: 11 May 2006
 Last Updated on STN: 31 May 2006
 Entered Medline: 30 May 2006
- L16 ANSWER 3 OF 42 HCAPLUS COPYRIGHT 2006 ACS on STN
- AN 2005:902982 HCAPLUS
- DN 143:242968
- TI Identification of mutation of paraoxonase 1 locus associated with predisposition to lacunar strokes and small vessel occlusion, and diagnostic and drug screening use
- IN Jannes, Jim; Hamilton-Bruce, Monica Anne; Koblar, Simon
- PA The Queen Elizabeth Hospital Research Foundation Inc., Australia; Adelaide Research & Innovation Pty. Ltd.
- SO PCT Int. Appl., 77 pp.
 - CODEN: PIXXD2
- DT Patent
- LA English
- FAN.CNT 1

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PATENT NO.
                        KIND
                               DATE
                                           APPLICATION NO.
                                                                  DATE
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                              20050825 WO 2005-AU218
    WO 2005078080
                        A1
                                                                  20050218
PΙ
        W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH,
             CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD,
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             RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML,
             MR, NE, SN, TD, TG
PRAI AU 2004-900790
                         Α
                               20040218
             THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT 2
             ALL CITATIONS AVAILABLE IN THE RE FORMAT
    ANSWER 4 OF 42 SCISEARCH COPYRIGHT (c) 2006 The Thomson Corporation on
L16
                                                       DUPLICATE 1
     2005:508032 SCISEARCH
AN
    The Genuine Article (R) Number: 905ZS
GA
    Structure/function analyses of rationally designed human serum
TI
    paraoxonase (HuPON1) mutants
ΑU
     Yeung D T (Reprint); Josse D; Lenz D E; Cerasoli D M
    USA, Med Res Inst Chem Def, Aberdeen Proving Ground, MD 21010 USA; Univ
CS
    Maryland, Baltimore, MD 21201 USA; Ctr Rech Serv Sante Armees, Dept
     Toxicol, F-38702 La Tronche, France
CYA
    USA; France
    FASEB JOURNAL, (7 MAR 2005) Vol. 19, No. 5, Part 2, Supp. [S], pp.
SO
    A1079-A1079.
     ISSN: 0892-6638.
    FEDERATION AMER SOC EXP BIOL, 9650 ROCKVILLE PIKE, BETHESDA, MD 20814-3998
PB
DT
    Conference; Journal
    English
LA
REC
    Reference Count: 0
    Entered STN: 26 May 2005
ED
    Last Updated on STN: 1 Dec 2005
    ANSWER 5 OF 42 HCAPLUS COPYRIGHT 2006 ACS on STN
L16
    2005:237537 HCAPLUS
AN
DN
    143:91740
TI
    Allelic variants of the human scavenger receptor class B type 1 and
    paraoxonase 1 on coronary heart disease: genotype-phenotype correlations
    Rodriguez-Esparragon, Francisco; Rodriguez-Perez, Jose C.;
ΑU
    Hernandez-Trujillo, Yaride; Macias-Reyes, Antonio; Medina, Alfonso;
    Caballero, Araceli; Ferrario, Carlos M.
    Research Unit, Nephrology and Cardiology Services, Hospital Universitario
CS
    de Gran Canaria Dr. Negrin, Las Palmas de Gran Canaria, Spain
    Arteriosclerosis, Thrombosis, and Vascular Biology (2005), 25(4), 854-860
SO
    CODEN: ATVBFA; ISSN: 1079-5642
PB
    Lippincott Williams & Wilkins
DT
    Journal
    English
LA
             THERE ARE 34 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT 34
             ALL CITATIONS AVAILABLE IN THE RE FORMAT
L16
    ANSWER 6 OF 42 HCAPLUS COPYRIGHT 2006 ACS on STN
AN
    2005:1110220 HCAPLUS
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cardiovascular disease in the Diabetes Heart Study

AU Burdon, K. P.; Langefeld, C. D.; Beck, S. R.; Wagenknecht, L. E.; Carr, J. J.; Freedman, B. I.; Herrington, D.; Bowden, D. W.

Association of genes of lipid metabolism with measures of subclinical

144:148032

DN

ΤI

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Cent. for Human Genomics, Wake Forest Univ. Sch. of Med., Winston-Salem,
CS
     NC, 27157, USA
     Journal of Medical Genetics (2005), 42(9), 720-724
SO
     CODEN: JMDGAE; ISSN: 0022-2593
     BMJ Publishing Group
PB
     Journal
DT
     English
LA
              THERE ARE 32 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT 32
              ALL CITATIONS AVAILABLE IN THE RE FORMAT
       ANSWER 7 OF 42
                         NTIS COPYRIGHT 2006 NTIS on STN
L16
                        NTIS Order Number: ADA443915/XAB
       2006(14):00902
AN
       Structure/Function Analyses of Human Serum Paraoxonase
TI
       (HuPON1) Mutants Designed from a DFPase-Like Homology Model.
       Journal article.
       Reprint: Structure/Function Analyses of Human Serum
       Paraoxonase (HuPON1) Mutants Designed from a
       DFPase-Like Homology Model.
ΑU
       Yeung, D. T.; Josse, D.; Nicholson, J. D.; Khanal, A.; McAndrew, C. W.
       Army Medical Research Inst. of Chemical Defense, Aberdeen Proving
CS
       Ground, MD. (075924000 396453)
NR
       ADA443915/XAB; USAMRICD-P04-019
       12p; 23 Aug 2004
DT
       Report
CY
       United States
LΑ
       English
       Pub. in Biochimica et Biophysica Acta, v1702 p67-77, 2004. Prepared in
NTE
       cooperation with Centre de Recherches du Service de Sante des Armees, La
       Tronche, France, and with University of Delaware, Newark, DE. The
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- os GRA&I0614
- ANSWER 8 OF 42 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on STN L16 AN
- 2004:283014 BIOSIS PREV200400283528 DN
- TТ DNA molecule encoding a variant paraoxonase and uses thereof.
- Salonen, Jukka T. [Inventor, Reprint Author]; Marchesani, Marja ΑU [Inventor]; Tuomainen, Tomi-Pekka [Inventor]; Kaikkonen, Jari [Inventor]
- CS Jannevirta, Finland ASSIGNEE: Oy Jurilab Ltd., Kuopio, Finland
- PΙ US 6740746 20040525
- SO Official Gazette of the United States Patent and Trademark Office Patents, (May 25 2004) Vol. 1282, No. 4. http://www.uspto.gov/web/menu/patdata.html e-file.

ISSN: 0098-1133 (ISSN print).

- DTPatent
- LA English
- ED Entered STN: 9 Jun 2004 Last Updated on STN: 9 Jun 2004
- ANSWER 9 OF 42 HCAPLUS COPYRIGHT 2006 ACS on STN L16
- 2004:756890 HCAPLUS AN
- DN 141:256526
- Variants of mammalian paraoxonases with increased solubility in bacterial TI expression hosts and near-normal kinetics and their selection, preparation, and therapeutic use
- Tawfik, Dan S.; Aharoni, Amir; Gaydukov, Leonid; Sussman, Joel L.; Silman, IN
- Yeda Research and Development Co. Ltd., Israel PA
- PCT Int. Appl., 240 pp. SO

CODEN: PIXXD2 DTPatent LA English FAN.CNT 1 PATENT NO. KIND DATE APPLICATION NO. DATE --------------______ ------A2 PΙ WO 2004078991 20040916 WO 2004-IL216 20040304 _ A3 WO 2004078991 20060105 W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM EP 1599598 A2 20051130 EP 2004-717208 20040304 AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK PRAI US 2003-451267P P 20030304 US 2003-512925P Р 20031022 WO 2004-IL216 W 20040304 ANSWER 10 OF 42 SCISEARCH COPYRIGHT (c) 2006 The Thomson Corporation on L16 STN **DUPLICATE 2** ΑN 2004:717190 SCISEARCH GΑ The Genuine Article (R) Number: 823UP TIStructure/function analyses of human paraoxonase-1 mutants Yeung D (Reprint); Josse D; Kirby S; Duryea C; Lenz D; Cerasoli D AU CS USA, Med Res Inst Chem Def, Aberdeen Proving Ground, MD 21010 USA; Serv Sante Armees, Ctr Rech, Toulon, France CYA USA; France FASEB JOURNAL, (14 MAY 2004) Vol. 18, No. 8, Supp. [S], pp. C144-C144. SO ISSN: 0892-6638. PΒ FEDERATION AMER SOC EXP BIOL, 9650 ROCKVILLE PIKE, BETHESDA, MD 20814-3998 DT Conference; Journal English LA REC Reference Count: 0 Entered STN: 3 Sep 2004 EDLast Updated on STN: 6 Oct 2005 => d 11-20 L16 ANSWER 11 OF 42 HCAPLUS COPYRIGHT 2006 ACS on STN 2004:78314 HCAPLUS AN DN 140:179620 ΤI Identification of paraoxonase 3 gene (PON3) missense mutations in a population of southern Italy AU Campo, Salvatore; Sardo, Adriana M.; Campo, Giuseppe M.; Avenoso, Angela; Castaldo, Maria; D'Ascola, Angela; Giunta, Elena; Calatroni, Alberto; Saitta, Antonino CS School of Medicine, Dept. Biochem., Physiological and Nutritional Sciences, University of Messina, Messina, 98125, Italy SO Mutation Research (2004), 546(1-2), 75-80 CODEN: MUREAV; ISSN: 0027-5107 PB Elsevier Science B.V. DTJournal

THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD

English

RE.CNT 12

LA

ALL CITATIONS AVAILABLE IN THE RE FORMAT

- L16 ANSWER 12 OF 42 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on STN
- AN 2004:235651 BIOSIS
- DN PREV200400236176
- TI PON1 M/L55 mutation protects high-risk patients against coronary artery disease.
- AU Oliveira, Simone A.; Mansur, Antonio P. [Reprint Author]; Ribeiro, Cristina C.; Ramires, Jose Antonio F.; Annichino-Bizzacchi, Joyce M.
- CS Medical School, Heart Institute (InCor), University of Sao Paulo, Av. Dr. Eneas Carvalho de Aguiar 44, CEP 05403-000, Sao Paulo, SP, Brazil corantonio@incor.usp.br
- SO International Journal of Cardiology, (March 2004) Vol. 94, No. 1, pp. 73-77. print.

 CODEN: IJCDD5. ISSN: 0167-5273.
- DT Article
- LA English
- ED Entered STN: 28 Apr 2004 Last Updated on STN: 28 Apr 2004
- L16 ANSWER 13 OF 42 MEDLINE on STN DUPLICATE 3
- AN 2004482197 MEDLINE
- DN PubMed ID: 15450851
- TI Structure/function analyses of human serum paraoxonase (HuPON1) mutants designed from a DFPase-like homology model.
- AU Yeung David T; Josse Denis; Nicholson James D; Khanal Akhil; McAndrew Christopher W; Bahnson Brian J; Lenz David E; Cerasoli Douglas M
- CS Biochemical Pharmacology Branch, Pharmacology Division, US Army Medical Research Institute of Chemical Defense, Aberdeen Proving Ground, MD 21010-5400, USA.
- NC 5P20RR015588 (NCRR)
- SO Biochimica et biophysica acta, (2004 Oct 1) Vol. 1702, No. 1, pp. 67-77. Journal code: 0217513. ISSN: 0006-3002.
- CY Netherlands
- DT Journal; Article; (JOURNAL ARTICLE)
- LA English
- FS Priority Journals
- EM 200411
- ED Entered STN: 29 Sep 2004
 Last Updated on STN: 2 Nov 2004
 Entered Medline: 1 Nov 2004
- L16 ANSWER 14 OF 42 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on STN
- AN 2003:292537 BIOSIS
- DN PREV200300292537
- TI New paraoxonase 1 polymorphism I102V and the risk of prostate cancer in Finnish men.
- AU Marchesani, Marja; Hakkarainen, Anna; Tuomainen, Tomi-Pekka; Kaikkonen, Jari; Pukkala, Eero; Uimari, Pekka; Seppala, Eija; Matikainen, Mika; Kallioniemi, Olli-P.; Schleutker, Johanna; Lehtimaki, Terho; Salonen, Jukka T. [Reprint Author]
- CS Research Institute of Public Health, University of Kuopio, 70211, Box 1627, Kuopio, Finland jukka.salonen@uku.fi
- Journal of the National Cancer Institute (Bethesda), (June 4 2003) Vol. 95, No. 11, pp. 812-818. print. ISSN: 0027-8874 (ISSN print).
- DT Article
- LA English
- ED Entered STN: 25 Jun 2003 Last Updated on STN: 25 Jun 2003
- L16 ANSWER 15 OF 42 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on

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- AN 2004:50093 BIOSIS
- DN PREV200400049274
- TI Paraoxonase 2 Ala148Gly mutation protects postmenopausal women from coronary artery disease.
- AU Mansur, A. P. [Reprint Author]; Oliveira, S. A.; Avakian, S. D. [Reprint Author]; Cesar, L. A. M. [Reprint Author]; Ramires, J. A. F. [Reprint Author]; Annichino-Bizzacchi, J. M. [Reprint Author]
- CS Medical School, Heart Institute (InCor), University of Sao Paulo, Sao Paulo, Brazil
- SO European Heart Journal, (August-September 2003) Vol. 24, No. Abstract Supplement, pp. 433. print.

 Meeting Info.: Congress of the European Society of Cardiology. Vienna, Austria. August 30-September 03, 2003. European Society of Cardiology. ISSN: 0195-668X (ISSN print).
- DT Conference; (Meeting)
 Conference; (Meeting Poster)
 Conference; Abstract; (Meeting Abstract)
- LA English
- ED Entered STN: 21 Jan 2004 Last Updated on STN: 21 Jan 2004
- L16 ANSWER 16 OF 42 HCAPLUS COPYRIGHT 2006 ACS on STN
- AN 2003:441718 HCAPLUS
- DN 139:208516
- TI Novel paraoxonase (PON1) nonsense and missense mutations predicted by functional genomic assay of PON1 status
- AU Jarvik, Gail P.; Jampsa, Rachel; Richter, Rebecca J.; Carlson, Chris S.; Rieder, Mark J.; Nickerson, Deborah A.; Furlong, Clement E.
- CS Departments of Medicine, The University of Washington, Seattle, WA, USA
- SO Pharmacogenetics (2003), 13(5), 291-295 CODEN: PHMCEE; ISSN: 0960-314X
- PB Lippincott Williams & Wilkins
- DT Journal
- LA English
- RE.CNT 13 THERE ARE 13 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT
- L16 ANSWER 17 OF 42 HCAPLUS COPYRIGHT 2006 ACS on STN
- AN 2004:335883 HCAPLUS
- DN 141:409279
- TI Association of paraoxonase 2 gene C311S variant with ischemic stroke in Chinese type 2 diabetes mellitus patients
- AU Wang, Xiaoyi; Xue, Yaoming; Wen, Shujuan; Zhang, Ningling; Ji, Zhong; Pan, Suyue
- CS Department of Endocrinology, General Military Hospital of Guangzhou, Guangzhou, Guangdong Province, 510010, Peop. Rep. China
- SO Zhonghua Yixue Yichuanxue Zazhi (2003), 20(3), 215-219 CODEN: ZYXZER; ISSN: 1003-9406
- PB Huaxi Yike Daxue
- DT Journal
- LA Chinese
- L16 ANSWER 18 OF 42 HCAPLUS COPYRIGHT 2006 ACS on STN
- AN 2003:267887 HCAPLUS
- DN 139:114895
- TI R192Q paraoxonase gene variant is associated with a change in HDL-cholesterol level during dietary caloric restriction in nondiabetic healthy males
- AU Obata, Tatsuo; Ito, Toshimitsu; Yonemura, Atsushi; Ayaori, Makoto; Nakamura, Haruo; Ohsuzu, Fumitaka
- CS First Department of Internal Medicine, National Defense Medical College, Saitama, Japan
- SO Journal of Atherosclerosis and Thrombosis (2003), 10(1), 57-62 CODEN: JATHEH; ISSN: 1340-3478

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PB
     Japan Atherosclerosis Society
DT
     Journal
     English
LА
              THERE ARE 29 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT 29
              ALL CITATIONS AVAILABLE IN THE RE FORMAT
L16 ANSWER 19 OF 42 HCAPLUS COPYRIGHT 2006 ACS on STN
     2003:389535 HCAPLUS
AN
DN
     139:160476
     Rapid genotyping of paraoxonase 55 and 192 mutations by melting point
ΤI
     analysis using real time PCR technology
     Pocsai, Zsuzsa; Toth, Zsuzsa; Paragh, Gyorgy; Szeles, Gyorgy; Adany, Roza
ΑU
CS
     Medical and Health Science Centre, School of Public Health, Department of
     Preventive Medicine, University of Debrecen, Debrecen, H-4012, Hung.
     Clinica Chimica Acta (2003), 332(1-2), 31-36
SO
     CODEN: CCATAR; ISSN: 0009-8981
PΒ
     Elsevier Science Ltd.
DT
     Journal
     English
LA
RE.CNT 10
              THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS RECORD
              ALL CITATIONS AVAILABLE IN THE RE FORMAT
      ANSWER 20 OF 42 BIOTECHDS COPYRIGHT 2006 THE THOMSON CORP. on STN
L16
      DUPLICATE 4
AN
      2003-03188 BIOTECHDS
      New genetic variants of human paraoxonase 1
ΤI
      (PON1) gene with polymorphisms, useful for treating disorders associated
      with PON1 isogene activity e.g. atherosclerosis or diabetes, or for
      screening drugs for treating these diseases;
         vector-mediated recombinant protein gene transfer and expression in
         host cell for use in drug screening and gene therapy
      ANASTASIO A E; CHEW A; CHOI J Y; DENTON R R; NANDABALAN K; PARKS K E;
AU
      STEPHENS J C
      GENAISSANCE PHARM INC
PΑ
      WO 2002066680 29 Aug 2002
ΑI
      WO 2001-US46896 6 Dec 2001
     WO 2001-5126 16 Feb 2001; WO 2001-5126 16 Feb 2001
PRAI
DT
      Patent
LA
      English
OS
      WPI: 2002-682769 [73]
=> d 21-30
L16 ANSWER 21 OF 42 HCAPLUS COPYRIGHT 2006 ACS on STN
AN
     2002:736050 HCAPLUS
DN
     137:275023
     Single nucleotide polymorphism (Ile101Val) in human paraoxonase gene
TI
     associated with diseases and diagnostic and therapeutic uses thereof
     Salonen, Jukka T.; Marchesani, Marja; Tuomainen, Tomi-Pekka; Kaikkonen,
IN
     Jari
PΑ
     Oy Jurilab Ltd., Finland
     PCT Int. Appl., 30 pp.
SO
     CODEN: PIXXD2
DT
     Patent
    English
LA
FAN.CNT 1
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                                          APPLICATION NO.
     PATENT NO.
                                                                 DATE
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     WO 2002074230
                                20020926
                                           WO 2002-FI231
                                                                   20020320
PΙ
                         A2
                               20030403
     WO 2002074230
                        A3
         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
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             GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
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LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,

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PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ,
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             GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA,
             GN, GQ, GW, ML, MR, NE, SN, TD, TG
                                            US 2001-811673
     US 2003003453
                          Α1
                                20030102
                                                                    20010320
                                20040525
     US 6740746
                          B2
                                            EP 2002-708387
                                                                    20020320
                          A2
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     EP 1370647
             AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
                          A1
                                20040722
                                            US 2003-691562
                                                                    20031024
     US 2004142349
PRAI US 2001-811673
                          Α
                                20010320
                                20020320
     WO 2002-FI231
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     2002:579314 HCAPLUS
DN
     137:323441
TI
     Relation between butyrylcholinesterase K variant, paraoxonase 1 (PON1) Q
     and R and apolipoprotein E &4 genes in early-onset coronary artery
     Nassar, Bassam A.; Darvesh, Sultan; Bevin, Lisa D.; Rockwood, Kenneth;
ΑU
     Kirkland, Susan A.; O'Neill, Blair J.; Bata, Iqbal R.; Johnstone, David
     E.; Title, Lawrence M.
CS
     Department of Pathology, Dalhousie University, Halifax, NS, Can.
     Clinical Biochemistry (2002), 35(3), 205-209
so
     CODEN: CLBIAS; ISSN: 0009-9120
PB
     Elsevier Science Inc.
DT
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LA
     English
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       2001(16):02587
ΤI
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       Sep 1997-21 Dec 2000.
ΑU
       Lockridge, 0.
       Nebraska Univ. Medical Center, Omaha. (034938000 417693)
CS
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       ADA387204/XAB
       315p; Jan 2001
NC
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DT
       Report
CY
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T.A
       English
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       Port Royal Road, Springfield, VA, 22161, USA.
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AN
     2001:190409 BIOSIS
     PREV200100190409
DN
ΤI
     PON2 gene variants are associated with clinical manifestations of
     cardiovascular disease in familial hypercholesterolemia patients.
     Leus, Frank R. [Reprint author]; Zwart, Miriam; Kastelein, John J. P.;
ΑU
     Voorbij, Hieronymus A. M.
     Department of Clinical Chemistry, University Hospital Utrecht,
CS
     Heidelberglaan 100, Rm. G03.647, 3584 CX, Utrecht, Netherlands
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f.leus@jc.azu.nl

- SO Atherosclerosis, (15 February, 2001) Vol. 154, No. 3, pp. 641-649. print. CODEN: ATHSBL. ISSN: 0021-9150.
- DT Article
- LA English
- ED Entered STN: 20 Apr 2001 Last Updated on STN: 18 Feb 2002
- L16 ANSWER 25 OF 42 HCAPLUS COPYRIGHT 2006 ACS on STN
- AN 2001:406986 HCAPLUS
- DN 135:193744
- TI Relationship of age-related myocardial infarction risk and Gln/Arg 192
 variants of the human paraoxonase-1 gene: the
 REGICOR study
- AU Senti, M.; Tomas, M.; Vila, J.; Marrugat, J.; Elosua, R.; Sala, J.; Masia, R.
- CS Institut Municipal d'Investigacio Medica, Lipids and Cardiovascular Epidemiology Unit, IMIM and Universitat Pompeu Fabra, Barcelona, E-08003, Spain
- SO Atherosclerosis (Shannon, Ireland) (2001), 156(2), 443-449 CODEN: ATHSBL; ISSN: 0021-9150
- PB Elsevier Science Ireland Ltd.
- DT Journal
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- RE.CNT 34 THERE ARE 34 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT
- L16 ANSWER 26 OF 42 HCAPLUS COPYRIGHT 2006 ACS on STN
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- DN 135:135684
- TI Polymorphism of apoprotein E (APOE), methylenetetrahydrofolate reductase (MTHFR) and paraoxonase (PON1) genes in patients with cerebrovascular disease
- AU Topic, Elizabeta; Simundic, Ana Maria; Stefanovic, Mario; Demarin, Vida; Vukovic, Vlasta; Lovrencic-Huzjan, Arijana; Zuntar, Irena
- CS Clinical Institute of Chemistry, School of Medicine, University of Zagreb and Sestre milosrdnice University Hospital, Zagreb, Zagreb, Croatia
- SO Clinical Chemistry and Laboratory Medicine (2001), 39(4), 346-350 CODEN: CCLMFW; ISSN: 1434-6621
- PB Walter de Gruyter GmbH & Co. KG
- DT Journal
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- RE.CNT 32 THERE ARE 32 CITED REFERENCES AVAILABLE FOR THIS RECORD
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- L16 ANSWER 27 OF 42 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on STN
- AN 2002:288744 BIOSIS
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- TI A new polymorphism in the PON1 gene predicts cardiovascular deaths: A prospective population-based study.
- AU Salonen, Jukka T. T. [Reprint author]; Marchesani, Maria; Tuomainen, Tomi-Pekka; Hakkarainen, Anna O.; Kaikkonen, Jari
- CS Research Inst of Public Health, Univ of Kuopio, Kuopio, Finland
- SO Circulation, (October 23, 2001) Vol. 104, No. 17 Supplement, pp. II.808. print.
 - Meeting Info.: Scientific Sessions 2001 of the American Heart Association. Anaheim, California, USA. November 11-14, 2001. American Heart Association.
 - CODEN: CIRCAZ. ISSN: 0009-7322.
- DT Conference; (Meeting)
 - Conference; Abstract; (Meeting Abstract)
- LA English
- ED Entered STN: 15 May 2002
 - Last Updated on STN: 15 May 2002

L16 ANSWER 28 OF 42 MEDLINE on STN DUPLICATE 5

AN 2000290007 MEDLINE

DN PubMed ID: 10831161

TI Differential effects of smoking on myocardial infarction risk according to the Gln/Arg 192 variants of the human paraoxonase gene.

AU Senti M; Aubo C; Tomas M

CS Lipids and Cardiovascular Epidemiology Unit, Institut Municipal d'Investigacio Medica, Barcelona, Spain.

SO Metabolism: clinical and experimental, (2000 May) Vol. 49, No. 5, pp. 557-9.

Journal code: 0375267. ISSN: 0026-0495.

CY United States

DT Journal; Article; (JOURNAL ARTICLE)

LA English

FS Priority Journals

EM 200006

ED Entered STN: 16 Jun 2000
Last Updated on STN: 16 Jun 2000
Entered Medline: 8 Jun 2000

L16 ANSWER 29 OF 42 MEDLINE on STN DUPLICATE 6

AN 2000189855 MEDLINE

DN PubMed ID: 10723110

TI Effect of physical activity on lipid levels in a population-based sample of men with and without the Arg192 variant of the human paraoxonase gene.

AU Senti M; Aubo C; Elosua R; Sala J; Tomas M; Marrugat J

CS Unitat de Lipids i Epidemiologia Cardiovascular, Institut Municipal d'Investigacio Medica, IMIM, Barcelona, Spain. msenti@imim.es

SO Genetic epidemiology, (2000 Mar) Vol. 18, No. 3, pp. 276-86. Journal code: 8411723. ISSN: 0741-0395.

CY United States

DT Journal; Article; (JOURNAL ARTICLE)

LA English

FS Priority Journals

EM 200005

ED Entered STN: 18 May 2000 Last Updated on STN: 18 May 2000 Entered Medline: 10 May 2000

L16 ANSWER 30 OF 42 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on STN

AN 2000:175266 BIOSIS

DN PREV200000175266

TI Common paraoxonase gene variants, mortality risk and fatal cardiovascular events in elderly subjects.

AU Heijmans, Bastiaan T.; Westendorp, Rudi G. J.; Lagaay, A. Margot; Knook, Dick L.; Kluft, Cornelis; Slagboom, P. Eline [Reprint author]

CS Gaubius Laboratory, TNO Prevention and Health, Zernikedreef 9, 2333 CK, Leiden, Netherlands

SO Atherosclerosis, (March, 2000) Vol. 149, No. 1, pp. 91-97. print. CODEN: ATHSBL. ISSN: 0021-9150.

DT Article

LA English

ED Entered STN: 3 May 2000 Last Updated on STN: 4 Jan 2002

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AN 2001:151271 BIOSIS

DN PREV200100151271

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     that are stable and soluble without amphiphilic additives.
     Original Title: Ingenierie de la paraoxonase humaine (PON1): Conception et
     construction de mutants solubles en l'absence de molecules amphiphiles.
     Borges, F. [Reprint author]; Viguie, N. [Reprint author]; Nachon, F.
ΑU
     [Reprint author]; Josse, D. [Reprint author]; Masson, P. [Reprint author]
     C.R.S.S.A., La Tronche, Grenoble, France
CS
     Travaux Scientifiques des Chercheurs du Service de Sante des Armees,
SO
     (2000) No. 21, pp. 37-38. print.
     ISSN: 0243-7473.
DT
     Article
LA
     French
ED
     Entered STN: 28 Mar 2001
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L16
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AN
     2000097604
                    MEDLINE
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     PubMed ID: 10634138
TI
     Mutations in the human paraoxonase 1 gene:
     frequencies, allelic linkages, and association with coronary artery
     Cascorbi I; Laule M; Mrozikiewicz P M; Mrozikiewicz A; Andel C; Baumann G;
AU
     Roots I; Stangl K
     Institute of Clinical Pharmacology, University Clinic Charite, Humboldt
CS
     University, Berlin, Germany.. ingolf.cascorbi@charite.de
     Pharmacogenetics, (1999 Dec) Vol. 9, No. 6, pp. 755-61.
SO
     Journal code: 9211735. ISSN: 0960-314X.
CY
     ENGLAND: United Kingdom
     Journal; Article; (JOURNAL ARTICLE)
DT
LΑ
     English
FS
     Priority Journals
     200001
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     Entered STN: 9 Feb 2000
ED
     Last Updated on STN: 9 Feb 2000
     Entered Medline: 28 Jan 2000
    ANSWER 33 OF 42 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on
L16
     STN
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AN
     PREV199900471499
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TI
     Mutations in the human paraoxonase gene:
     Frequencies, allelic linkages and association with coronary artery
     disease.
     Stangl, K. [Reprint author]; Cascorbi, I.; Laule, M. [Reprint author];
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     Stangl, V. [Reprint author]; Baumann, G. [Reprint author]; Roots, I.
     Med. Klinik, Kardiologie, Charite, Humboldt Universitaet, Campus Mitte,
CS
     Berlin, Germany
     European Heart Journal, (Aug., 1999) Vol. 20, No. ABSTR. SUPPL., pp. 178.
SO
     Meeting Info.: XXIst Congress of the European Society of Cardiology.
     Barcelona, Spain. August 28-September 1, 1999. European Society of
     Cardiology.
     CODEN: EHJODF. ISSN: 0195-668X.
DT
     Conference; (Meeting)
     Conference; Abstract; (Meeting Abstract)
     Conference; (Meeting Poster)
LA
     English
     Entered STN: 9 Nov 1999
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     Last Updated on STN: 9 Nov 1999
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     1999:384202 HCAPLUS
     131:180577
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     Paraoxonase 1 Mutations in a Turkish Population
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Nacak, Muradiye; Tapanyigit, E. Ebru; Roots, Ivar

- Institute of Clinical Pharmacology, University Clinic Charite, Humboldt CS University, Berlin, Germany
- Toxicology and Applied Pharmacology (1999), 157(3), 174-177 SO CODEN: TXAPA9; ISSN: 0041-008X
- Academic Press PB
- Journal DT
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- 1998:440131 HCAPLUS AN
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- Kao, Yan-Lin; Donaghue, Kim; Chan, Albert; Knight, John; Silink, Martin AII
- Ray Williams Institute of Paediatric Endocrinology, Diabetes and CS Metabolism, The Royal Alexandra Hospital for Children, Westmead, 2145, Australia
- Journal of Clinical Endocrinology and Metabolism (1998), 83(7), 2589-2592 SO CODEN: JCEMAZ; ISSN: 0021-972X
- PB Endocrine Society
- DTJournal
- English LA
- THERE ARE 20 CITED REFERENCES AVAILABLE FOR THIS RECORD RE.CNT 20 ALL CITATIONS AVAILABLE IN THE RE FORMAT
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- DN PubMed ID: 9672215
- Molecular variant of the human paraoxonase TI /arylesterase gene is associated with central retinal vein occlusion in the Japanese population.
- Murata M; Nakagawa M; Takahashi S ΑU
- Department of Ophthalmology, Yamagata University School of Medicine, CS Japan.
- Ophthalmologica. Journal international d'ophtalmologie. International SO journal of ophthalmology. Zeitschrift fur Augenheilkunde, (1998) Vol. 212, No. 4, pp. 257-9.
 - Journal code: 0054655. ISSN: 0030-3755.
- CY Switzerland
- Journal; Article; (JOURNAL ARTICLE) DT
- LΑ English
- Priority Journals FS
- EΜ 199809
- Entered STN: 6 Oct 1998 ED Last Updated on STN: 6 Oct 1998 Entered Medline: 21 Sep 1998
- L16 ANSWER 37 OF 42 HCAPLUS COPYRIGHT 2006 ACS on STN
- 1998:164903 HCAPLUS AN
- 128:307033 DN
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- Sanghera, Dharambir K.; Saha, Nilmani; Kamboh, M. Ilyas ΑU
- 130 DeSoto Street, Graduate School of Public Health, Department of Human CS Genetics, University of Pittsburgh, Pittsburgh, PA, 15261, USA
- Atherosclerosis (Shannon, Ireland) (1998), 136(2), 217-223 so CODEN: ATHSBL; ISSN: 0021-9150
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DN
     127:118118
     Genetic mapping and gene structure of mouse paraoxonase/arylesterase and
TI
     site-directed mutagenesis of human paraoxonase
     /arylesterase
AU
     Sorenson, Robert Christian
     Univ. of Michigan, Ann Arbor, MI, USA
CS
     (1997) 152 pp. Avail.: UMI, Order No. DA9722097
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     From: Diss. Abstr. Int., B 1997, 58(2), 520
     Dissertation
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     English
LΑ
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     PubMed ID: 9437206
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     A 192Arg variant of the human paraoxonase
     (HUMPONA) gene polymorphism is associated with an increased risk for
     coronary artery disease in the Japanese.
     Zama T; Murata M; Matsubara Y; Kawano K; Aoki N; Yoshino H; Watanabe G;
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     Ishikawa K; Ikeda Y
    Department of Medicine, School of Medicine, Keio University, Tokyo, Japan.
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    Arteriosclerosis, thrombosis, and vascular biology, (1997 Dec) Vol. 17,
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    No. 12, pp. 3565-9.
     Journal code: 9505803. ISSN: 1079-5642.
CY
    United States
     Journal; Article; (JOURNAL ARTICLE)
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     English
     Priority Journals
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     199802
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     Entered STN: 17 Feb 1998
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     Entered Medline: 3 Feb 1998
L16 ANSWER 40 OF 42 SCISEARCH COPYRIGHT (c) 2006 The Thomson Corporation on
     STN
     1997:626141 SCISEARCH
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     The Genuine Article (R) Number: XE898
GA
     An (192) Arg variant of the human paraoxonase
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     (HUMPONA) gene polymorphisms is associated for with the increased risk for
     coronary artery disease in the Japanese
     Zama T (Reprint); Murata M; Matsubara Y; Kawano K; Aoki N; Yoshino H;
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     Watanabe G; Ishikawa K; Ikeda Y
     KEIO UNIV, SCH MED, DEPT MED, TOKYO 160, JAPAN; KYORIN UNIV, DEPT MED 2,
CS
     TOKYO, JAPAN; HIBIYA MED CTR, SAKURA BANK, TOKYO, JAPAN
CYA JAPAN
     THROMBOSIS AND HAEMOSTASIS, (JUN 1997) Supp. [S], pp. P2129-P2129.
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     ISSN: 0340-6245.
     F K SCHATTAUER VERLAG GMBH, P O BOX 10 45 45, LENZHALDE 3, D-70040
PB
     STUTTGART, GERMANY.
     Conference; Journal
DT
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     LIFE
     English
LA
REC Reference Count: 0
     Entered STN: 1997
ED
     Last Updated on STN: 1997
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L16 ANSWER 41 OF 42 MEDLINE on STN

DUPLICATE 10

AN 96292331 MEDLINE

DN PubMed ID: 8675673

TI A variant of human paraoxonase/arylesterase

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(HUMPONA) gene is a risk factor for coronary artery disease.
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     Serrato M; Marian A J
     Division of Cardiology, Department of Medicine, Baylor College of
CS
     Medicine, Houston, Texas 77030, USA.
     The Journal of clinical investigation, (1995 Dec) Vol. 96, No. 6, pp.
SO
     3005-8.
     Journal code: 7802877. ISSN: 0021-9738.
CY
     United States
     Journal; Article; (JOURNAL ARTICLE)
DT
LA
     Abridged Index Medicus Journals; Priority Journals
FS
     199608
EM
     Entered STN: 22 Aug 1996
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     Last Updated on STN: 22 Aug 1996
     Entered Medline: 13 Aug 1996
    ANSWER 42 OF 42 SCISEARCH COPYRIGHT (c) 2006 The Thomson Corporation on
L16
     STN
                                                         DUPLICATE 11
     1992:152692 SCISEARCH
AN
GA
     The Genuine Article (R) Number: HG719
     A POINT MUTATION APPEARS TO ACCOUNT FOR THE HUMAN
ΤI
     SERUM PARAOXONASE ARYLESTERASE POLYMORPHISM
     ADKINS S (Reprint); GAN K; MODY M; LADU B
ΑU
     UNIV MICHIGAN, SCH PUBL HLTH, DEPT ENVIRONM & IND HLTH, ANN ARBOR, MI
CS
     48109; UNIV MICHIGAN, SCH MED, DEPT PHARMACOL, ANN ARBOR, MI 48109; UNIV
     MICHIGAN, SCH MED, DEPT ANESTHESIOL, ANN ARBOR, MI 48109
CYA
    USA
     FASEB JOURNAL, (26 FEB 1992) Vol. 6, No. 4, Part 1, pp. A1582-A1582.
SO
     ISSN: 0892-6638.
     FEDERATION AMER SOC EXP BIOL, 9650 ROCKVILLE PIKE, BETHESDA, MD
PΒ
     20814-3998.
DT
     Conference; Journal
     LIFE
FS
     English
LA
REC Reference Count: 2
     Entered STN: 1994
ED
     Last Updated on STN: 1994
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L3
              5 DUP REM L2 (22 DUPLICATES REMOVED)
          13181 S JOLY, ?/AU
L4
              0 S L1 AND L4
L5
              8 S L4 AND LUCIFERASE AND GFP
L6
              4 DUP REM L6 (4 DUPLICATES REMOVED)
L7
           6625 S · PARAOXONASE
L8
           4548 S L8 AND HUMAN
L9
            947 S L9 AND (VARIANT OR MUTA? OR SUBSTITUTION)
L10
           2115 S L8 (10A) HUMAN
L11
            104 S L11(10A) (VARIANT OR MUTA? OR SUBSTITUTION)
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L14
           1896 S L8 (5A) HUMAN
             78 S L14(5A) (VARIANT OR MUTA? OR SUBSTITUTION)
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ENTRY

SESSION

FULL ESTIMATED COST 115.16 115.37

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